

Evaluation of Recent Updates to the Spectroscopy of CO₂ and CH₄ in the Thermal Infrared Using Observations from TES and IASI

M. J. Alvarado¹ (malvarad@aer.com), V. H. Payne², K. E. Cady-Pereira¹, S. Kulawik², A. P. Chase^{1,*}, E. Mlawer¹, J.-L. Moncet¹, M. W. Shephard³, and J. Delamere^{1,**}

¹Atmospheric and Environmental Research (AER) ²Jet Propulsion Laboratory, California Institute of Technology ³Environment Canada *Now at University of Maine **Now at Tech-X

1. Introduction

The accuracy of retrieved profiles of greenhouse gases such as CO₂ and CH₄ depends on the accuracy of the radiative transfer model used in the retrieval. Uncertainties in spectroscopic line parameters and continua are the primary limitations on the accuracy of molecular absorption in radiative transfer models, and so reducing these uncertainties is critical to ensuring future scientific progress. As part of our work with the TES instrument, AER regularly examines and validates potential spectroscopic improvements for inclusion in the forward model of the TES retrieval algorithm.

Here we present the results of several validation studies of recent updates to the spectroscopic parameters for CO₂ and CH₄ in AER's line-by-line radiative transfer model LBLRTM against measurements from the Tropospheric Emission Spectrometer (TES) made during the HIAPER Pole-to-Pole Observations (HIPPO) of Carbon Cycle and Greenhouse Gases Study and a global dataset of near-nadir measurements from the Infrared Atmospheric Sounding Instrument (IASI). We focus on the spectral residuals in the main thermal infrared bands of CH₄ and CO₂.

2. LBLRTM is an accurate and flexible radiative transfer model that is the basis of the forward model for a number of satellite programs, including TES and IASI.

LBLRTM v12.1 with AER v3.1 line parameters (released November 2011, rtweb.aer.com)

Spectroscopy based on HITRAN 2008 (Rothman et al., 2009) along with:

- H₂O
 - Line positions and intensities (10-2500 cm⁻¹): Coudert et al. (2008).
 - Air-broadened half-widths, temp. dep. and pressure shifts (350-667 cm⁻¹): Delamere et al. (2010).
- CO₂
 - Lamouroux et al. (2010) first order line coupling parameters (P-, Q-, and R-branches).
 - Line intensities and positions (597-2500 cm⁻¹) from the Carbon Dioxide Spectral Database (CDSD) (Tashkun et al., 2003; Flaud et al., 2003).
- CH₄
 - HITRAN 2008 with first-order line coupling for ν₄ and ν₃ bands (Tran et al., 2006).
- MT_CKD v2.6.2 Continuum
 - Updates to CO₂ and self-broadened H₂O in the 2400 cm⁻¹ region (Mlawer et al., 2012).

LBLRTM v11.3 with TES v1.4 line parameters (released November 2007)

Spectroscopy based on HITRAN 2000 (Rothman et al., 2003) along with:

- H₂O from HITRAN 2006 updates.
- CO₂, P-, Q-, and R-branch line coupling based on Niro et al. (2005).
- CH₄ (922.65-1678.33 cm⁻¹) and CO supplied by Linda Brown of JPL.
- O₃ from Wagner et al., 2002.
- MT_CKD v2.0 Continuum.

LBLRTM v9.4+ with AER v1.0 line parameters (released January 2005)

As in LBLRTM v11.3 with TES v1.4 except CO₂ Q-branch line coupling only and MT_CKD v1.2 Continuum.

3. Methods

TES is a Fourier Transform Spectrometer (FTS) aboard the NASA Aura polar orbiting satellite. Spectral range 650–1325 cm⁻¹ and 1900–2250 cm⁻¹, 0.06 cm⁻¹ resolution, footprint of 5.3×8.3 km.

IASI is a FTS aboard the MetOp-A polar orbiting satellite. Spectral range 645–2760 cm⁻¹, 0.5 cm⁻¹ resolution, footprint of ~18 km, swath of ~2400 km.

a) TES-HIPPO forward model (FM) study

We use TES spectra measured during the HIPPO I campaign (Jan. 2009) that are nearly coincident with the aircraft. We use the observed aircraft profiles of CO₂ and CH₄ (supplied by the HIPPO science team, S. Wofsy, lead PI) and examine the forward model residuals using different line parameters.

b) TES-HIPPO retrieval study

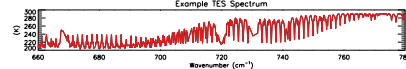
We use spectra measured by TES during the HIPPO I and II campaigns. We performed retrievals of temperature, H₂O, CO₂, HDO, N₂O, CH₄, cloud OD, and surface emissivity using different line parameters.

c) IASI retrieval study

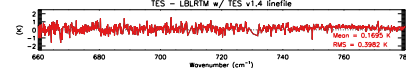
We use a subset of the spectra from Matricardi, 2009. These are clear-sky, nighttime, ocean profiles. We performed retrievals of temperature, H₂O, O₃, CO, and CH₄. A priori profiles are from ECMWF model or TES climatology

4. Preliminary TES-HIPPO FM Results: CO₂

Example TES
Spectrum



Mean Residuals
LBLv11.3 w/TES v1.4



Mean Residuals
LBL v12.1 w/AER v3.1

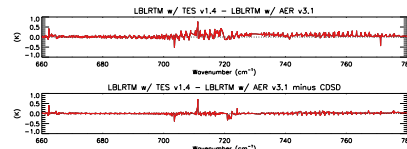


Mean Residuals
LBLv12.1 w/AER v3.1
Minus CDSD



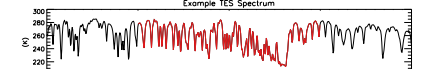
Using the CDSD positions and strengths included in LBLRTM v12.1 (and AER v3.1) increases the RMS of the mean forward model residuals in the CO₂ ν₂ band.

Left: Mean residuals for 6 scans over ocean with cloud OD < 0.1.
Below: Mean differences between LBL runs using the TES v1.4, AER v3.1, and AER v3.1 minus CDSD line parameters.

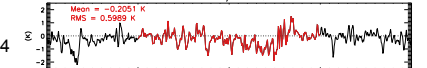


5. Preliminary TES-HIPPO FM Results: CH₄ and N₂O

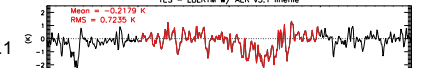
Example TES
Spectrum



Mean Residuals
LBL v11.3 w/ TES v1.4

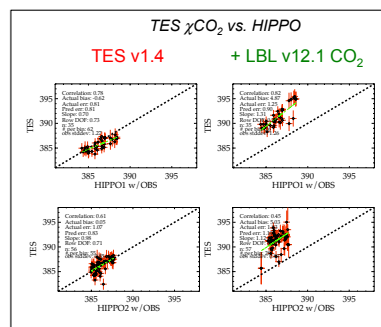
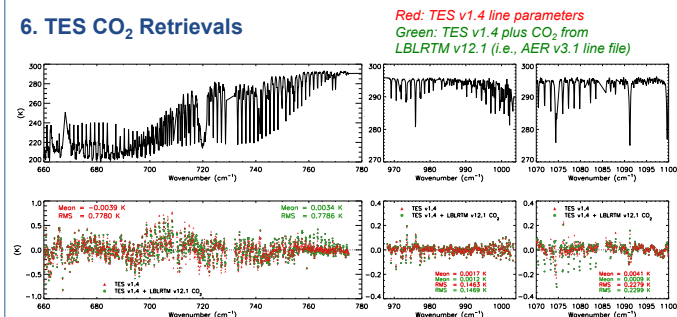


Mean Residuals
LBL v12.1 w/ AER v3.1

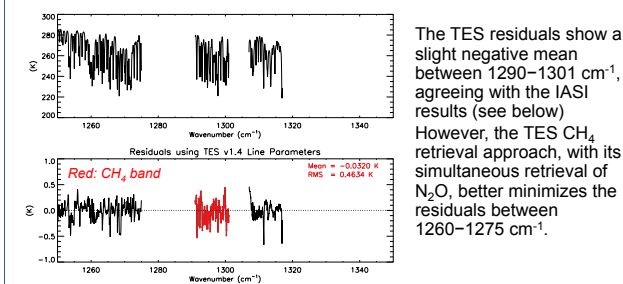


The updated CH₄ and N₂O spectroscopic parameters included in LBLRTM v12.1 don't appear to improve the forward model residuals in this spectral range.

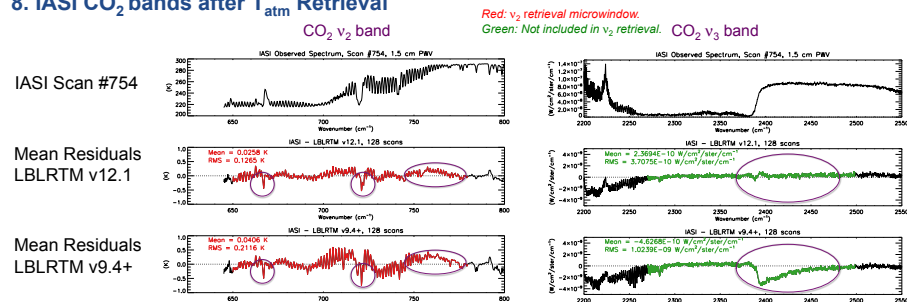
6. TES CO₂ Retrievals



7. TES CH₄ Retrievals

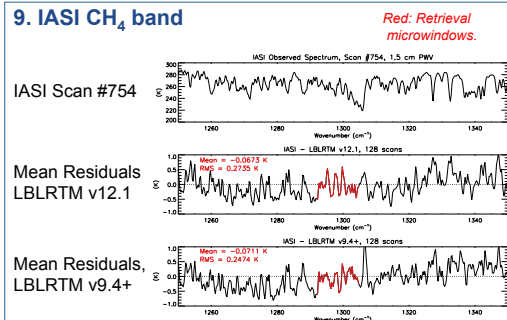


8. IASI CO₂ bands after T_{atm} Retrieval



In LBLRTM v9.4+, there were significant residuals near the v₃ bandhead after the v₂ band temperature retrieval. Recent updates to the MT_CKD continuum included in LBLRTM v12.1 have dramatically improved the model performance in this region.

9. IASI CH₄ band



It is not clear if the CH₄ spectroscopic updates have improved the residuals. While the mean residual is now closer to 0, the RMS is larger.

The retrieved CH₄ profiles (not shown) have a clear high bias, suggesting that spectroscopic errors remain in this region.

10. Conclusions

The LBLRTM v12.1 CO₂ spectroscopy is remarkably consistent between the CO₂ v₂ and v₃ bands. However, the performance of the TES CO₂ retrieval, which does not use the CO₂ v₃ band, is degraded by adding the CDSD positions and strengths in LBLRTM v12.1 to the TES line parameters.

The LBLRTM v12.1 CH₄ spectroscopy is not a clear improvement over that in the TES v1.4 line parameters or LBLRTM v9.4+. Strong residual features remain between 1295–1301 cm⁻¹, suggesting further spectroscopic work is needed.

References

- Blumstein et al., *Proc. SPIE* 6684, 66840H, 2007.
- Coudert et al., *J. Mol. Spectrosc.*, 251, 339, 2008.
- Delamere et al., *J. Geophys. Res.*, 115, D17106, 2010.
- Filaud et al., *ESA TN-LPM-FAC-02*, 2003.
- Hoke et al., *In: IRS 88: Current Problems in Atm. Rad.*, 368-371, 1989.
- Lamoureaux et al., *JQSRT*, 111, 2321, 2010.
- Maselli et al., *Atmos. Chem. Phys.*, 11, 1009-1021, 2011.
- Matricardi, *Atmos. Chem. Phys.*, 9, 6889-6913, 2009.
- Milaver et al., *Phil. Trans. Roy. Soc.*, 370, 2520-2556, 2012.
- Rodgers and Connor, *J. Geophys. Res.*, 108, 4116, 2003.
- Rothman et al., *JQSRT*, 82, 5-44, 2003.
- Rothman et al., *JQSRT*, 110, 533-572, 2009.
- Strow et al., *JQSRT*, 52, 281-294, 1994.
- Tashkun et al., 14th Int. Symp. on High Res. Mol. Spectrosc., 2003.
- Wagner et al., *J. Geophys. Res.*, 107, 4626, 2002.